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We claim:

A peptide of the formula: R¹ - X¹ - X² - R²
 wherein X¹ is an aromatic amino acid residue;
 X² is any amino acid residue;
 R¹ is NH₂ - or an amino acid sequence X³ - X⁴ - X⁵

wherein X³ is an aliphatic amino acid residue having a side chain hydroxyl group and X⁴ and X⁵ are the same or different and are any amino acid residue and wherein R² is a sequence of 1 to 3 amino acid residues which are the same or different and are aliphatic amino acid residues.

3. The peptide of claim 1

wherein R¹ is NH₂-;

X¹ is an aromatic amino acid;

X² is Glu or Ala and

R² is Gly, Gly-Gly, Gly-Gly-Gly or

sarcosine.

- 4. The peptide of claim 3 wherein X^1 is Phe and X^2 is Glu.
- 5. The peptide of claim 2 having an amino acid sequence selected from the group

consisting of:

- (a) Thr-Asp-Ile-Phe-Glu-Gly-Gly (Sequence ID NO:8);
- (b) Thr-Ala-Ile-Phe-Glu-Gly-Gly (Sequence ID NO:3);
- (c) Thr-Asp-Ala-Phe-Glu-Gly-Gly (Sequence ID NO:4);
 and

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- (d) Thr-Asp-Ile-Phe-Ala-Gly-Gly (Sequence ID NO:6).
- 6. The peptide of claim 3 having an amino acid sequence selected from the group

consisting of:

- (a) Phe-Glu-Gly-Gly-Gly (Sequence ID NO:9);
- (b) Phe-Glu-Gly; and
- (c) Phe-Glu-Sarcosine.
- 7. The peptide of claim 1 wherein R² is a sequence of 1 to 3 amino acid residues which are the same or different and are selected from the group consisting of glycine, sarcosine, azetidine, nipecotic acid and pipecotic acid.
- 8. The peptide of claim 3 wherein R² is a sequence of 1 to 3 amino acid residues which are the same or different and are selected from the group consisting of glycine, sarcosine, azetidine, nipecotic acid and pipecotic acid.
- 9. The peptide of any of claims 1 to 8 wherein at least one amino acid is a D amino acid.
 - 10. The peptide of claim 4 or 6 wherein Phe and Glu are D amino acids.
- 25 11. A peptide having the amino acid sequence Ser-Gly-Glu-Gly-Val-Arg (Sequence ID NO:1).
- 12. A pharmaceutical composition comprising a peptide of any of claims 1 to 11 and a pharmaceutically30 acceptable carrier.
 - 13. A method for treating or preventing SIRS-induced hypotension in a mammal comprising administering to the mammal an effective amount of a peptide of any of claims
- 1, 2, 5, 7, 9 or 11 or of an effective fragment or derivative of said peptide.
 - 14. A method for treating or preventing anaphylactic

hypotension in a mammal comprising administering to the mammal an effective amount of a peptide of any of claims 1 to 10, or of an effective fragment or derivative of said peptide.

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- 15. A method of reducing or preventing an anaphylactic reaction in a mammal comprising administering an effective amount of a peptide of any of claims 1 to 10 or of an effective fragment or derivative of said peptide to the mammal.
- 16. A method of reducing or preventing an endotoxic reaction in a mammal comprising administering an effective amount of a peptide of any of claims 1, 2, 5,
- 15 7, 9 or 11 or an effective fragment or derivative of said peptide to the mammal.
- 17. A method for treating an inflammatory disorder in a mammal comprising administering to the mammal an
 20 effective amount of a peptide of any claims 1, 2, 5, 7, 9 or 10 or of an effective fragment or derivative of the peptide to the mammal.
- 18. The method of claim 17 wherein the inflammatory
 25 disorder is selected from the group consisting of a
 rheumatic disorder, inflammatory bowel disease, postischemic inflammation or systemic inflammatory response
 syndrome.
- 30 19. An antibody which specifically recognises an epitope of a peptide of any of claims 1 to 11.
- 20. A method of determining the peptide SGP-T or the peptide SGPS in a biological fluid comprising obtaining a sample of the biological fluid and determining the peptide in the fluid by immunoassay employing an antibody of claim 19.